

GenCore version 5.1.5
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OM nucleic - nucleic search, using sw model
Run on: June 1, 2003, 17:59:15 ; Search time 497 seconds
(without alignments)
10113.602 Million cell updates/sec

Title: US-09-625-573-1
Perfect score: 2232
Sequence: 1 GGATTGAACAGGACGATT.....TATACTATCTGTGATAAAG 2232

Scoring table: OLIGO_NUC
Gapop 60.0, Gapext 60.0

Searched: 2185239 seqs, 1125999159 residues

Word size : 0
Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : N_Geneseq_101002: *
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3: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1982.DAT: *
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21: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2000.DAT: *
22: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT: *
23: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT: *
24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	2232	100.0	2232	16	AAQ96297 Human monocyte che
2	1152	51.6	143068	21	AAF21105 Human low adenosin
3	1152	51.6	143068	21	AAF21272 Human low adenosin
4	1152	51.6	143068	21	AAA34983 Human adenosine re
5	1152	51.6	143068	21	AAA35150 Human adenosine re
6	1152	51.6	143068	21	ABL68124 Ovary cancer relat
7	1152	51.6	149412	21	ABL68124 Human adenosine re
8	1152	51.6	152740	21	AAF21273 Human low adenosin
9	980	43.9	1979	16	AAQ96298 Human monocyte che

10	941	42.2	1083	22	AAAS12140	Human wild-type CC
11	890	39.9	1083	22	AAAS12139	Human CCR2-64I pol
12	839	37.6	1083	18	AAAT96976	Human monocyte che
13	839	37.6	1083	23	AB197976	Non-endogenous hum
14	65	2.9	461	20	AAV84136	HIV-1 co-receptor
15	65	2.9	792	18	AAAT90116	cDNA for inactive
16	65	2.9	1056	22	AAAD13198	Human G-protein ch
17	65	2.9	1056	22	AAAD13299	Human G-protein ch
18	65	2.9	1056	24	ABK51870	DNA encoding human
19	65	2.9	1059	19	AAV23992	Human CC-CKR5 codi
20	65	2.9	1059	23	AB197978	Non-endogenous hum
21	65	2.9	1071	24	ABA97319	Human chemokine (C
22	65	2.9	1071	20	AAV84125	HIV-1 co-receptor
23	65	2.9	1225	19	AAAT76920	DNA encoding human
24	65	2.9	1225	24	ABA02317	Human CC chemokine
25	65	2.9	1225	24	ABA02318	Human CCR5 Gln 55
26	65	2.9	1255	19	AAAT76919	DNA encoding human
27	65	2.9	1344	20	AAV84159	HIV-1 co-receptor
28	65	2.9	1376	22	AAH26903	Human HIV-1 co-rec
29	65	2.9	1414	18	AAAT44042	Human G-protein ch
30	65	2.9	1414	21	AAZ91481	Human G-protein ch
31	65	2.9	1414	22	AAAD13181	Human HDGRL0 cDNA
32	65	2.9	1414	22	AAAD13282	DNA encoding human
33	65	2.9	1414	22	AAAF26390	cDNA for inactive
34	65	2.9	1414	22	AAAF26390	cDNA for human CCR
35	65	2.9	1414	24	ABK51853	Human CCR5 cDNA se
36	65	2.9	1442	18	AAAT90118	Human chemokine re
37	65	2.9	1477	18	AAAT90117	Human chemokine re
38	65	2.9	1477	22	AAAF87099	Human chemokine re
39	65	2.9	1557	18	AAAT99542	Human low adenosin
40	65	2.9	3383	18	AAAT85161	Human adenosine re
41	65	2.9	3383	21	AAAF21271	Human chemokine re
42	65	2.9	3383	21	AAA35149	Human chemokine re
43	65	2.9	3383	22	AAAD08577	Human chemokine re
44	65	2.9	5674	20	AAZ24738	Human chemokine (C
45	65	2.9	9141	24	ABA97318	Human chemokine (C

ALIGNMENTS

RESULT 1
ID AAQ96297
ID AAQ96297 standard; cDNA; 2232 BP.
XX AC AAQ96297;
XX DT 29-DEC-1995 (first entry)
XX DE Human monocyte chemoattractant protein-1 receptor MCP-1RA.
XX DE Monocyte chemoattractant protein-1 receptor; MCR-1R; chemokine; ss.
XX KW Homo sapiens.
XX OS Homo sapiens.
XX FH Key Location/Qualifiers
XX FT CDS 40...1161
XX FT /*tag= a
XX PN WO9519436-A.
XX PD 20-JUL-1995.
XX PF 11-JAN-1995; 95WO-US00476.
XX PR 13-JAN-1994; 94US-0182962.
XX PA (REGC) UNIV CALIFORNIA.
XX PI Charo I, Coughlin S;
XX DR WPI; 1995-263866/34.
XX DR P-PSDB; AAR79165.

xx DNA encoding monocyte chemo-attractant protein-1 receptor - used partic.
pt for identifying antagonists and for treating diseases characterised by
pt monocyte infiltrates

Disclosure; Fig 1; 84pp; English.

To identify and clone new members of the chemokine receptor gene family, degenerate oligo primers were designed corresp. to the conserved sequences R79167 in the second and R79168 in the third transmembrane domains of the MIP-1alpha/RANTES receptor, the IL-8 receptors and the HUMSTRS orphan receptor (Genbank Accession #M99293). The degenerate oligo incorporating EcoRI and XhoI sites at their 5' ends are Q96299 and Q96300. Amplification of cDNA derived from MM6 cells with the primers yielded a number of PCR products. One cDNA appeared to encode a novel protein. To obtain a full-length version of this clone, a MM6 cDNA library was constructed in pFRG and probed with the PCR product. A 2.1 kb cDNA clone was obtained. Analysis of additional clones in the MM6 cDNA library revealed a second sequence that was identical to the 2.1 kb cDNA sequence first obtained from the 5' UTR through the putative seventh transmembrane domain but contained a different cytoplasmic tail. The second sequence appears to represent alternative splicing of the carboxyl-terminal tail of the MCP-1R protein. The two sequences are denoted MCP-1RA and MCP-1RB (see Q96297/R79165 & Q96298/R79166). Active mature MCP-1RA has a predicted mol. wt. of about 42,000 daltons. MCP-1RB has a mol. wt. of about 41,000 daltons.

Sequence 2232 BP; 602 A; 464 C; 508 G; 658 T; 0 other;

Query Match	100.0%	Score 2232;	DB 16;	Length 2232;
Best Local Similarity	100.0%;	Pred. No. 0;		
Matches 2232;	Conservative	0;	Mismatches	0;
			Indels	0;
			Gaps	0;

1	GGATTGAACAAGGAGCGATTTCCCCAGTACATCCACAACATGCTGCCACATCTCGTTCT	60
1	GGATTGAACAAGGAGCGATTTCCCGAGTACATCCACAACATGCTGCCACATCTCGTTCT	60
61	CGGTTTATCAGAATATACCAAGAGGCGGTGAAGAAGTACCACCTTTTGGTTATGAT	120
61	CGGTTTATCAGAATATACCAAGAGGCGGTGAAGAAGTACCACCTTTTGGTTATGAT	120
121	TACGGTGTCCCTGTGCATAAATTTGAGTGAAGCAAAATTTGGGGCCCAACTCTGCGCTCG	180
121	TACGGTGTCCCTGTGCATAAATTTGAGTGAAGCAAAATTTGGGGCCCAACTCTGCGCTCG	180
181	CNCTACTCGCTGGTGTCTACCTTTGGTTTTGTGGCAACATGCTGCTGCTCCTCATCTTA	240
181	CNCTACTCGCTGGTGTCTACCTTTGGTTTTGTGGCAACATGCTGCTGCTCCTCATCTTA	240
241	ATAAACTGCAAAAAGCTGAAGTCTTGACTGACATTTTACTGCTCAACTGGGCCATCTCT	300
241	ATAAACTGCAAAAAGCTGAAGTCTTGACTGACATTTTACTGCTCAACTGGGCCATCTCT	300
301	GATCGCTTTTCTTATTAATCTCCCATTTGGGGTCTACTCTGCTGCAAAAGAGTGGGTC	360
301	GATCGCTTTTCTTATTAATCTCCCATTTGGGGTCTACTCTGCTGCAAAAGAGTGGGTC	360
361	TTTGGGAATGCAATGTGCAAAATTTATCAAGGGCTGTATCAATCGGTTATTTTGGCGGA	420
361	TTTGGGAATGCAATGTGCAAAATTTATCAAGGGCTGTATCAATCGGTTATTTTGGCGGA	420
421	ATCTTCTTCATCATCTCCTGCAATCGATAGATACCTGGCTATTGTCCATGCTGTGTT	480
421	ATCTTCTTCATCATCTCCTGCAATCGATAGATACCTGGCTATTGTCCATGCTGTGTT	480
481	GCCTTAAAGCCAGGAGGTCACTTTGGGGTGGTGACAAGTGTGATCACTGGTGGTG	540
481	GCCTTAAAGCCAGGAGGTCACTTTGGGGTGGTGACAAGTGTGATCACTGGTGGTG	540
541	GCTGTGTTTGTCTCTGCCAGGAATCATCTTTACTAAATGCCAAGAAGATCTGT	600
541	GCTGTGTTTGTCTCTGCCAGGAATCATCTTTACTAAATGCCAAGAAGATCTGT	600

1681 CTTCTAGGCTTGTGCTCCAAAGACCTTTTCTGTTTCTGTTTCTGTTATCATATGAGTCATGC 1740
Db CTTCTAGGCTTGTGCTCCAAAGACCTTTTCTGTTTCTGTTTCTGTTATCATATGAGTCATGC 1740
1741 GTTTAATACATTCGAGTGTTCAGTGTCTTCCAGATGTCCTTGTATGCTCATATTTGTC 1800
Db GTTTAATACATTCGAGTGTTCAGTGTCTTCCAGATGTCCTTGTATGCTCATATTTGTC 1800
1801 CTAATTTGCCAGTGGGAATCTTAATCAAAATTCGCTTCTTAATCAAAAGCTTTTAAACCT 1860
Db CTAATTTGCCAGTGGGAATCTTAATCAAAATTCGCTTCTTAATCAAAAGCTTTTAAACCT 1860
1861 ATTGGTAAGAATGGAGGTGGAGAGCTCCCTGAAGTAAGCAAGAACTTTCTCTTAGT 1920
Db ATTGGTAAGAATGGAGGTGGAGAGCTCCCTGAAGTAAGCAAGAACTTTCTCTTAGT 1920
1921 CGAGCCAAAGTAAAGATGTTCTTATGTTGCCAGTGTGTTCTGATCTGATGCAAGCAAG 1980
Db CGAGCCAAAGTAAAGATGTTCTTATGTTGCCAGTGTGTTCTGATCTGATGCAAGCAAG 1980
1981 AAACACTGGGCTTCTAGAACCCAGCAACTTGGGAACCTAGACTCCCAAGCTGGACTATGGC 2040
Db AAACACTGGGCTTCTAGAACCCAGCAACTTGGGAACCTAGACTCCCAAGCTGGACTATGGC 2040
2041 TCTACTTTTCAGCCACATGGCTTAAGAGGTTTCAGAAAGAGTGGGGACAGAGCAAGAC 2100
Db TCTACTTTTCAGCCACATGGCTTAAGAGGTTTCAGAAAGAGTGGGGACAGAGCAAGAC 2100
2101 TTTACCTTCATATATTTGATGTCCTTAATGAATGCAATAAATGTTAAGTGTGATGGA 2160
Db TTTACCTTCATATATTTGATGTCCTTAATGAATGCAATAAATGTTAAGTGTGATGGA 2160
2161 TGAATGTAAATACATGTTTAAACAACTATGATTTGGAAATTAATCAATGCTATAACTA 2220
Db TGAATGTAAATACATGTTTAAACAACTATGATTTGGAAATTAATCAATGCTATAACTA 2220
2221 TGTGATAAAG 2232
Db TGTGATAAAG 2232

RESULT 2
AAF21105
ID AAF21105 standard; DNA; 143068 BP.
XX AC AAF21105;
XX DT 14-MAR-2001 (first entry)

Human low adenosine antisense oligonucleotide related sequence #2672.

Low adenosine antisense oligonucleotide; phosphorothioate; allergy;
human; airway disorder; bronchoconstriction; lung inflammation;
surfactant depletion; respiratory; bronchodilator; antiinflammatory;
immunosuppressive; antiasthmatic; analgesic; hypotensive; cytostatic;
respiratory obstruction; pulmonary obstruction; impeded respiration;
surfactant hypoproduction; pulmonary vasoconstriction; asthma; RDS;
respiratory distress syndrome; emphysema; pain; cystic fibrosis; allergic rhinitis;
chronic obstructive pulmonary disease; pulmonary infection; bronchitis;
cancer; ss.

Homo sapiens.

W0200062736-A2.

26-OCT-2000.

24-MAR-2000; 2000WO-US08020.

06-APR-1999; 99US-0127958.

(UYEC-) UNIV EAST CAROLINA.

(NYCE/) NYCE J W.

Nyce JW;

WPI; 2000-679539/66.

Low adenosine (A) content antisense oligonucleotides which do not trigger adenosine receptors during metabolism, useful e.g. for treating cancers and respiratory obstructions -

Disclosure; Page 924-957; 1592pp; English.

The present invention describes low adenosine (A) content antisense oligonucleotides and compositions (I) comprising them. In the antisense oligonucleotides the A is replaced by a 'Universal' or alternative base. (I) can have respiratory, bronchodilator, antiinflammatory, analgesic, immunosuppressive, antiasthmatic, hypotensive and cytostatic activities. The antisense oligonucleotides and (I) can be used to down-regulate the expression and/or activity of target polypeptides associated with lung/respiratory disorders and malignancies, such as stimulating and activating peptide factors and transmitters, transcription factors, immunoglobulins and antibodies, antibody receptors, cytokines and chemokines, endogenously produced specific and non-specific enzymes, binding proteins, adenosine receptors, bradykinin receptors, central chemokine receptors, adenosine receptors, and non-nervous system receptors, CNS and peripheral nervous and non-nervous system peptide transmitters, defensins, growth factors, vasoactive peptides and receptors, binding proteins and malignancy associated proteins. The antisense oligonucleotides may be used in this way to treat disorders including respiratory obstruction (especially pulmonary obstruction and/or bronchoconstriction) and/or lung inflammation, allergy(ies) and/or surfactant hypoproduction which are associated with a disease or condition selected from pulmonary vasoconstriction, inflammation, allergies, asthma, impeded respiration, respiratory distress syndrome (RDS), pain, cystic fibrosis (CF), allergic rhinitis (AR), pulmonary hypertension, emphysema, chronic obstructive pulmonary disease (COPD), pulmonary transplantation rejection, pulmonary infections, bronchitis, and/or cancer. AAF18434 to AAF21543 represent human polynucleotide fragments and antisense oligonucleotides used in the exemplification of the present invention.

Sequence 143068 BP; 41194 A; 30122 C; 32403 G; 39349 T; 0 other;

Query Match 51.6%; Score 1152; DB 21; Length 143068;

Best Local Similarity 99.8%; Pred. No. 0;

Matches 1252; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 979 AGCCTTTTTCACATAGCTCTTGGCTGTAGGATTCGCCACCTCCAAACACCAAGTGTGGA 1038
Db 48253 AGCCTTTTTCACATAGCTCTTGGCTGTAGGATTCGCCACCTCCAAACACCAAGTGTGGA 48312
QY 1039 GGTCCAGGAGTGAGACAGGAAAGAAATGTGAAGTGACTACAGAGGACTCTCGATGCT 1098
Db 48313 GGTCCAGGAGTGAGACAGGAAAGAAATGTGAAGTGACTACAGAGGACTCTCGATGCT 48372
QY 1099 CGTGGAAAAGAAAGTCAATTTGGCAGAGCCCTGAAGCCAGTCTTCAGGACAAAGGA 1158
Db 48373 CGTGGAAAAGAAAGTCAATTTGGCAGAGCCCTGAAGCCAGTCTTCAGGACAAAGGA 48432
QY 1159 GCCTAGACAGAGAATGACAGATCTCTCTTTGGAAATACACAGTCTGGTTCACAGATG 1218
Db 48433 GCCTAGACAGAGAATGACAGATCTCTCTTTGGAAATACACAGTCTGGTTCACAGATG 48492
QY 1219 TGTGATTCACAGTGTGAATCTTGGTGTCTACGTTACAGGAGGAGGCTGAGAGGAGAG 1278
Db 48493 TGTGATTCACAGTGTGAATCTTGGTGTCTACGTTACAGGAGGAGGCTGAGAGGAGAG 48552
QY 1279 AGACTCCAGCTGGGTTGGAAAACAGTATTTCCAACTACCTTCCAGTTCTTCATTTTG 1338
Db 48553 AGACTCCAGCTGGGTTGGAAAACAGTATTTCCAACTACCTTCCAGTTCTTCATTTTG 48612
QY 1339 AATACAGGATAGAGTTTTCAGACTTTTTTAAATAGTAAATAATTAAGCTGAAAC 1398

Db 48613 AATACAGGATAGAGTTGAGACTTTTAAATAGTAAATAAATAAATAAAGCTGAAC 48672
QY 1399 TCGAAGCTTGAATGTAAGAGTGTAGTTGAGTTGCTATCATCTCAAGAGTGAAT 1458
Db 48673 TCGAAGCTTGAATGTAAGAGTGTAGTTGAGTTGCTATCATCTCAAGAGTGAAT 48732
QY 1459 GCTGTATTAGTCACAGAGATAATCTAGCTTTGAGCTTAAGAAATTTGAGCAGGTGGTAT 1518
Db 48733 GCTGTATTAGTCACAGAGATAATCTAGCTTTGAGCTTAAGAAATTTGAGCAGGTGGTAT 48792
QY 1519 GTTTGGGAGACTGCTGAGTCAACCAATAGTTGATGTCAGAGAGTTGGAAGTGTG 1578
Db 48793 GTTTGGGAGACTGCTGAGTCAACCAATAGTTGATGTCAGAGAGTTGGAAGTGTG 48852
QY 1579 ATCTGTGGGACATAGGCTTATGTCAGCATCTGAGCATCTAAGTAATGATGCTGTTGAATCA 1638
Db 48853 ATCTGTGGGACATAGGCTTATGTCAGCATCTGAGCATCTAAGTAATGATGCTGTTGAATCA 48912
QY 1639 CAGTATAGCTCCATCGCTGTCATCTCAGCTGGATCCCATCTCTCAGGCTTCTGCTGCA 1698
Db 48913 CAGTATAGCTCCATCGCTGTCATCTCAGCTGGATCCCATCTCTCAGGCTTCTGCTGCA 48972
QY 1699 AAGCCCTTTGTTGTTTGTATCATTAATGAGTCATGCTGTTAATCATCATTCAGT 1758
Db 48973 AAGCCCTTTGTTGTTTGTATCATTAATGAGTCATGCTGTTAATCATCATTCAGT 49032
QY 1759 GTTTCAGTGTGTCAGATGCTCTGATGCTCATATGTTCCCTAATTTCCAGTGGGAA 1818
Db 49033 GTTTCAGTGTGTCAGATGCTCTGATGCTCATATGTTCCCTAATTTCCAGTGGGAA 49092
QY 1819 CTCTTAATCAATGCTCTTAAATCAAGCTTTTAAACCTTATGTTAAAGATGAAG 1878
Db 49093 CTCTTAATCAATGCTCTTAAATCAAGCTTTTAAACCTTATGTTAAAGATGAAG 49152
QY 1879 GTGGAGAAGCTCCTCAAGTAAAGAGTCTTCTTATGTCGAGCCCAAGTTAAGAAATG 1938
Db 49153 GTGGAGAAGCTCCTCAAGTAAAGAGTCTTCTTATGTCGAGCCCAAGTTAAGAAATG 49212
QY 1939 TTCTTATGTCGAGTGTGTTTCTGATGCTGATGCAAGCAAGAACACACACACACACAC 1998
Db 49213 TTCTTATGTCGAGTGTGTTTCTGATGCTGATGCAAGCAAGAACACACACACACACAC 49272
QY 1999 ACCAGGCAACTGGGAAGTCTGAGTCCCAAGTGGAGTATGGCTCTACTTTCAGGCCACAT 2058
Db 49273 ACCAGGCAACTGGGAAGTCTGAGTCCCAAGTGGAGTATGGCTCTACTTTCAGGCCACAT 49332
QY 2059 GCCTAAGAGGTTTCAGAAAGAGTGGGGACAGAGCAGAGACTTTTCACCTTCATATATT 2118
Db 49333 GCCTAAGAGGTTTCAGAAAGAGTGGGGACAGAGCAGAGACTTTTCACCTTCATATATT 49392
QY 2119 GTATGATCCTTAATGATCAATAATGTTAAGTGGTGGTGAATGAAATGTAATACTGTT 2178
Db 49393 GTATGATCCTTAATGATCAATAATGTTAAGTGGTGGTGAATGAAATGTAATACTGTT 49452
QY 2179 TTTAACAACATGATTTGGAAATTAATCAATGCTATTAACATGTTGATATAAG 2332
Db 49453 TTTAACAACATGATTTGGAAATTAATCAATGCTATTAACATGTTGATATAAG 49506

RESULT 3

AAFP21272
ID AAF21272 standard; DNA; 143068 BP.
XX AAF21272;
AC AAF21272;
DT 14-MAR-2001 (first entry)
XX Human low adenosine antisense oligonucleotide related sequence #2839.
DE Low adenosine antisense oligonucleotide; phosphorothioate; allergy;
XX human; airway disorder; bronchoconstriction; lung inflammation;
KW surfactant depletion; respiratory; bronchodilator; antiinflammatory;
KW

immunosuppressive; antiasthmatic; analgesic; hypotensive; cytostatic;
respiratory obstruction; pulmonary obstruction; impeded respiration;
surfactant hypoproduction; pulmonary vasoconstriction; asthma; RDS;
respiratory distress syndrome; pain; cystic fibrosis; allergic rhinitis;
pulmonary hypertension; emphysema; pulmonary transplantation rejection;
chronic obstructive pulmonary disease; pulmonary infection; bronchitis;
cancer; ss.
Homo sapiens.
WO2000062736-A2.
26-OCT-2000.
24-MAR-2000; 2000WO-US08020.
06-APR-1999; 99US-0127958.
(UYEC-) UNIV EAST CAROLINA.
(NYCE/) NYCE J W.
Nyce JW;
WPI; 2000-679539/66.
Low adenosine (A) content antisense oligonucleotides which do not
trigger adenosine receptors during metabolism, useful e.g. for treating
cancers and respiratory obstructions -
Disclosure; Page 1186-1219; 1592pp; English.
The present invention describes low adenosine (A) content antisense
oligonucleotides and compositions (I) comprising them. In the antisense
oligonucleotides the A is replaced by a 'Universal' or alternative base.
(I) can have respiratory, bronchodilator, antiinflammatory, analgesic,
immunosuppressive, antiasthmatic, hypotensive and cytostatic activities.
The antisense oligonucleotides and (I) can be used to down-regulate the
expression and/or activity of target polypeptides associated with the
lung/respiratory disorders and malignancies, such as stimulating and
activating peptide factors and transmitters, such as stimulating and
immunoglobulins and antibodies, antibody receptors, cytokines and
chemokines, endogenously produced specific and non-specific enzymes,
binding proteins, adhesion molecules and their receptors, cytokine and
chemokine receptors, adenosine receptors, bradykinin receptors, central
nervous system (CNS) and peripheral nervous and non-nervous system
receptors, CNS and peripheral nervous and non-nervous system peptide
transmitters, defensins, growth factors, vasoactive peptides and
receptors, binding proteins and malignancy associated proteins. The
antisense oligonucleotides may be used in this way to treat disorders
including respiratory obstruction (especially pulmonary obstruction
and/or bronchoconstriction) and/or lung inflammation, allergy(ies)
and/or surfactant hypoproduction which are associated with a disease or
condition selected from pulmonary vasoconstriction, inflammation,
allergies, asthma, impeded respiration, respiratory distress syndrome
(RDS), pain, cystic fibrosis (CF), allergic rhinitis (AR), pulmonary
hypertension, emphysema, chronic obstructive pulmonary disease (COPD),
pulmonary transplantation rejection, pulmonary infections, bronchitis,
and/or cancer. AAF18434 to AAF21543 represent human polynucleotide
fragments and antisense oligonucleotides used in the exemplification of
the present invention.

Sequence 143068 BP; 41194 A; 30122 C; 32403 G; 39349 T; 0 other;
Query Match 51.6%; Score 1152; DB 21; Length 143068;
Best Local Similarity 99.8%; Pred. No. 0;
Matches 1252; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 979 AGCCTTTTTCACATAGCTCTTGGCTGTAGGATTCGCCCATCCAAAACAGTGTGTGGA 1038
|||||
Db 48253 AGCCTTTTTCACATAGCTCTTGGCTGTAGGATTCGCCCATCCAAAACAGTGTGTGGA 48312
|||||
QY 1039 GGTCCAGGAGTGAGACCAGGAAGAATGTGAAAGTGACTACACAGGACTCTTCGATGT 1098
|||||

Db 48313 GGTCCAGGAGTGAGACAGCAAGAAATGTAAGTGAATGACTACACAAGGACTCCTCGATGGT 48372
QY 1099 CBTGGAAGAGAAAGTCAATGTCGACAGCCCTGAGCCAGCTTTCAGGACAAAGAGGA 1158
Db 48373 CBTGGAAGAGAAAGTCAATGTCGACAGCCCTGAGCCAGCTTTCAGGACAAAGAGGA 48432
QY 1159 GCTAGACAGCAAGTACAGATCTCTGCTTGGAAATACACAGTCTGGCTTCACAGATG 1218
Db 48433 GCTAGACAGCAAGTACAGATCTCTGCTTGGAAATACACAGTCTGGCTTCACAGATG 48492
QY 1219 TGTGATTCACAGTGTGAATCTTGGTGTCTAGCTTACAGGACGAGGCTGAGAGGAG 1278
Db 48493 TGTGATTCACAGTGTGAATCTTGGTGTCTAGCTTACAGGACGAGGCTGAGAGGAG 48552
QY 1279 AGACTCAGCTGGTGGAAACAGTATTTCCAAACTACCTTCCAGTCTCCTCATTTTGG 1338
Db 48553 AGACTCAGCTGGTGGAAACAGTATTTCCAAACTACCTTCCAGTCTCCTCATTTTGG 48612
QY 1339 AATACAGGATAGAGTTCAGACTTTTAAATAGTAAATAAATAAAGCTGAAAC 1398
Db 48613 AATACAGGATAGAGTTCAGACTTTTAAATAGTAAATAAATAAAGCTGAAAC 48672
QY 1399 TGCACCTGTAAATGTGGTAAAGTGTAGTTGAGTTGCTATCATGTCAACGCTGAAAT 1458
Db 48673 TGCACCTGTAAATGTGGTAAAGTGTAGTTGAGTTGCTATCATGTCAACGCTGAAAT 48732
QY 1459 GCTGTATAGTACAGAGATAATCTAGCTTTGAGCTTAAAGATTTTTCAGGAGGTGGTAT 1518
Db 48733 GCTGTATAGTACAGAGATAATCTAGCTTTGAGCTTAAAGATTTTTCAGGAGGTGGTAT 48792
QY 1519 GTTTGGGAGACTGCTGAGTCAACCAATAGTTGTTGATGGCAGAGTGGAGTGTGTG 1578
Db 48793 GTTTGGGAGACTGCTGAGTCAACCAATAGTTGTTGATGGCAGAGTGGAGTGTGTG 48852
QY 1579 ATCTGTGGCAGATTTAGCTATGTCAGTCAGCATCTAAGTAATGATGCTGTTGAATCA 1638
Db 48853 ATCTGTGGCAGATTTAGCTATGTCAGTCAGCATCTAAGTAATGATGCTGTTGAATCA 48912
QY 1639 CAGTATAGCTCCATGCTGCTCATCTCAGCTGAGTCTCATTTCTCAGGCTTGTGCA 1698
Db 48913 CAGTATAGCTCCATGCTGCTCATCTCAGCTGAGTCTCATTTCTCAGGCTTGTGCA 48972
QY 1699 AAGGCTTTTGTGTTTGTATCATATGAGTCAATGCTGTTAAATCAATTCAGT 1758
Db 48973 AAGGCTTTTGTGTTTGTATCATATGAGTCAATGCTGTTAAATCAATTCAGT 49032
QY 1759 GTTTCAGTCTCGCAGATGCTCTGATGCTCATATGTTCCCTAATTTGCCAGTGGAA 1818
Db 49033 GTTTCAGTCTCGCAGATGCTCTGATGCTCATATGTTCCCTAATTTGCCAGTGGAA 49092
QY 1819 CTCTTAATCAATTTGGCTTCTAATCAAGCTTTTAAACCTATTTGTAAGAAATGAAG 1878
Db 49093 CTCTTAATCAATTTGGCTTCTAATCAAGCTTTTAAACCTATTTGTAAGAAATGAAG 49152
QY 1879 GTGAGAGCTCCTGAGTAAGCAAGACTTTCCTTCTAGTCAAGCAAGTAAAGAAATG 1938
Db 49153 GTGAGAGCTCCTGAGTAAGCAAGACTTTCCTTCTAGTCAAGCAAGTAAAGAAATG 49212
QY 1939 TTCTTATTTGCCAGTGTGTTCTGATGCTGATGCAAGCAAGAACTGGGCTTCTAGA 1998
Db 49213 TTCTTATTTGCCAGTGTGTTCTGATGCTGATGCAAGCAAGAACTGGGCTTCTAGA 49272
QY 1999 ACAGGCACTTGGAACTAGCTCCCAAGCTGGACTATGCTCTACTTTCAGGCCACAT 2058
Db 49273 ACAGGCACTTGGAACTAGCTCCCAAGCTGGACTATGCTCTACTTTCAGGCCACAT 49332
QY 2059 GGCTAAAGAGGTTTCAGAAAGAGTGGGAGCAGACAGACAGAACTTTCACCTTCATATTT 2118
Db 49333 GGCTAAAGAGGTTTCAGAAAGAGTGGGAGCAGACAGACAGAACTTTCACCTTCATATTT 49392
QY 2119 GTATGATCCTTAATGATGCAATAAATGTTAGTTGATGGTGAATGTAATGTAATGTT 2178
Db 49393 GTATGATCCTTAATGATGCAATAAATGTTAGTTGATGGTGAATGTAATGTAATGTT 49452

QY 2179 TTTTAACTATGATTTGGAAATAAATCAATGCTATCACTATGTTGATAAAG 2232
Db 49453 TTTTAACTATGATTTGGAAATAAATCAATGCTATCACTATGTTGATAAAG 49506

RESULT 4
AAA34983
ID AAA34983 standard; DNA; 143068 BP.
XX
AC AAA34983;
XX
XX 28-JUL-2000 (first entry)
XX
XX Human adenosine receptor related polynucleotide SEQ ID NO:2672.

Human; adenosine receptor; low adenosine antisense oligonucleotide; phosphorothioate; impaired respiration; inflammation; allergy; allergic disease; bronchoconstriction; inhibitor; antiinflammatory; antiallergic; antisthmatic; cytotatic; analgesic; impaired airway; lung disease; ischaemic condition; pulmonary vasoconstriction; asthma; respiratory distress syndrome; pain; cystic fibrosis; emphysema; pulmonary hypertension; chronic obstructive pulmonary disease; COPD; cancer; leukaemia; lymphoma; carcinoma; metastasis; ss.

Homo sapiens.
XX
XX WO200009525-A2.
XX
XX 24-FEB-2000.
XX
XX 03-AUG-1999; 99WO-US17712.
XX
XX 03-AUG-1998; 98US-0095212.
XX
XX (UYEC-) UNIV EAST CAROLINA.
XX
XX Nyce JW;
XX
XX WPI; 2000-205971/18.
XX
XX New antisense oligonucleotides useful for treating e.g. pulmonary vasoconstriction, inflammation, allergies, asthma, hypertension, bronchitis, emphysema, respiratory distress syndrome, ischemia or cancers -
XX
XX Disclosure; Page 851-882; 1343pp; English.

The present invention describes a new composition comprising an antisense oligonucleotide (ON) with low adenosine (up to 15%), which targets nucleic acids involved in bronchoconstriction, allergies, and/or inflammation. The ON can have antiinflammatory, antiallergic, antiasthmatic, cytotatic and analgesic activities. The compositions are useful for the treatment of diseases associated with inflammation, impaired airways, including lung disease and diseases whose secondary effects afflict the lungs of a subject. They can be used for treating e.g. ischaemic conditions, pulmonary vasoconstriction, allergies, asthma, impaired respiration, respiratory distress syndrome, pain, cystic fibrosis, pulmonary hypertension, emphysema, chronic obstructive pulmonary disease (COPD), and cancers which may metastasise to the lungs, including carcinomas, and cancers which may metastasise to the lungs, including breast and prostate cancer. The reduction of the adenosine content of the ONs reduces side effects. The A-containing ONs break down with the release of deoxyadenosine which activates adenosine receptors causing bronchoconstriction and inflammation. AAA32313 to AAA35312 represent the nucleotide sequences given in the sequence listing from the present invention, which correspond to SEQ ID NO:1 to 2815, and then the last 185 sequences are also called SEQ ID NO:1 to 185, and then the sequences differ from the previously named sequences. SEQ ID NO:11 to 1680 (AAA32323 to AAA33992) are specifically claimed ONs from the present invention. N.B. Sequences given in the disclosure of the present invention do not match up with their corresponding SEQ ID NO: sequences given in the sequence listing.

1. *Chlorophyll a* (Chl a) is the primary photosynthetic pigment in most plants and algae. It is a green pigment that absorbs light energy in the blue and red regions of the visible spectrum. Chl a is essential for the light-dependent reactions of photosynthesis, where it converts light energy into chemical energy in the form of ATP and NADPH.

PR	27-SEP-2000;	2000US-235720P;
PR	27-SEP-2000;	2000US-235840P;
PR	27-SEP-2000;	2000US-235863P;
PR	28-SEP-2000;	2000US-236028P;
PR	28-SEP-2000;	2000US-236032P;
PR	28-SEP-2000;	2000US-236033P;
PR	28-SEP-2000;	2000US-236034P;
PR	28-SEP-2000;	2000US-236310P;
PR	28-SEP-2000;	2000US-236111P;
PR	29-SEP-2000;	2000US-2356842P;
PR	29-SEP-2000;	2000US-236891P;
PR	02-OCT-2000;	2000US-237172P;
PR	02-OCT-2000;	2000US-237173P;
PR	02-OCT-2000;	2000US-237278P;
PR	02-OCT-2000;	2000US-237294P;
PR	02-OCT-2000;	2000US-237295P;
PR	03-OCT-2000;	2000US-237316P;
PR	03-OCT-2000;	2000US-237425P;
PR	03-OCT-2000;	2000US-237598P;
PR	03-OCT-2000;	2000US-237604P;
PR	03-OCT-2000;	2000US-237606P;
PR	03-OCT-2000;	2000US-237608P;
PR	01-NOV-2000;	2000US-244867P;
PR	01-NOV-2000;	2000US-245084P;

	Query Match	51.6%	Score 1152;	DB 24;	Length 143068;
	Best Local Similarity	99.8%;	Pred. No. 0;		
	Matches 1252;	Conservative 0;	Mismatches 2;	Indels 0;	Gaps 0;
QY	979	AGCCTTTTTCACATAGCTCTTGGCTGTAGATTGCCCCACTCCAAAAACCAAGTGTGTGGA	1038		
Db	48253	AGCCTTTTTCACATAGCTCTTGGCTGTAGATTGCCCCACTCCAAAAACCAAGTGTGTGGA	48312		
QY	1039	GGTCCAGGAGTGTAGACCAGGAAAGAATGTCAAAGTGTACTACACAAGGACTCTCTCGATGGT	1098		
Db	48313	GGTCCAGGAGTGTAGACCAGGAAAGAATGTCAAAGTGTACTACACAAGGACTCTCTCGATGGT	48372		
QY	1099	CGTGGAAAGGAAAGTCAATTGSCAGAGCCCTTGAGCCAGCTCTTCAGGACAAAGAAGGA	1158		
Db	48373	CGTGGAAAGGAAAGTCAATTGSCAGAGCCCTTGAGCCAGCTCTTCAGGACAAAGAAGGA	48432		

QY	1159	GCCTAGAGACAGAAATGACAGATCTCTGCTTTGGAAATCACACGCTCTGGCTTCACAGATG	1218
Db	48433	GCCTAGAGACAGAAATGACAGATCTCTGCTTTGGAAATCACACGCTCTGGCTTCACAGATG	48492
QY	1219	TGTCATTACACAGTGTGAATCTTGGTGCTCTACGTTACCAAGCAGGAAGGCTTCAGAGGAGAG	1278
Db	48493	TGTCATTACACAGTGTGAATCTTGGTGCTCTACGTTACCAAGCAGGAAGGCTTCAGAGGAGAG	48552
QY	1279	AGACTCCAGCTGGGTTGGAAACAGATATTTTCCAAACTACCTTCCAGTTCTCATTTTGT	1338
Db	48553	AGACTCCAGCTGGGTTGGAAACAGATATTTTCCAAACTACCTTCCAGTTCTCATTTTGT	48612
QY	1339	AATACAGCATAGAGTTCAGACTTTTTTAAATAGTAAAAATAAAATTAAGCTGAAAAAC	1398
Db	48613	AATACAGCATAGAGTTCAGACTTTTTTAAATAGTAAAAATAAAATTAAGCTGAAAAAC	48672
QY	1399	TGCAACTTGTAAATGTGGTAAAGAGTTAGTTTGAGTTGCTATCATGTCAAACGCTGAAAAAT	1458
Db	48673	TGCAACTTGTAAATGTGGTAAAGAGTTAGTTTGAGTTGCTATCATGTCAAACGCTGAAAAAT	48732
QY	1459	GCTGTATTAGTCACAGAGATAATTCTCTAGCTTTTGAGCTTTAAGAATTTTGACGAGTGGTAT	1518
Db	48733	GCTGTATTAGTCACAGAGATAATTCTCTAGCTTTTGAGCTTTAAGAATTTTGACGAGTGGTAT	48792
QY	1519	GTTTGGGAGACTGCTGAGTCAACCAATAGTTGTTCATTTGCGCAGGAGTTTGGAGTGTGTG	1578
Db	48793	GTTTGGGAGACTGCTGAGTCAACCAATAGTTGTTCATTTGCGCAGGAGTTTGGAGTGTGTG	48852
QY	1579	ATCTGTGGGCACATTAGCCTATGTGCATGCGACGACTCAAGTAATGATGCTTTTGAATCA	1638
Db	48853	ATCTGTGGGCACATTAGCCTATGTGCATGCGACGACTCAAGTAATGATGCTTTTGAATCA	48912
QY	1639	CAGTATACGCTCCATCGCTGTCATCTCAGCTTGGATCTCGATTCCTCAGGCTTGTGCGCA	1698
Db	48913	CAGTATACGCTCCATCGCTGTCATCTCAGCTTGGATCTCGATTCCTCAGGCTTGTGCGCA	48972
QY	1699	AAAGCCTTTTGTGTTTGTGTTTGTATCATTTAAGTTCATGCGTTTAAATCACATTCGAGT	1758
Db	48973	AAAGCCTTTTGTGTTTGTGTTTGTATCATTTAAGTTCATGCGTTTAAATCACATTCGAGT	49032
QY	1759	GTTTCAGTGTCTGCAGATGCTCTTGATGCTCATATTGTTCCCTAATTTGCCAGTGGGAA	1818
Db	49033	GTTTCAGTGTCTGCAGATGCTCTTGATGCTCATATTGTTCCCTAATTTGCCAGTGGGAA	49092
QY	1819	CTCCTAAATCAAATTTGGCTTCTAATCAAAGCTTTTAAACCCCTATTGGTGAAGAATGGAAG	1878
Db	49093	CTCCTAAATCAAATTTGGCTTCTAATCAAAGCTTTTAAACCCCTATTGGTGAAGAATGGAAG	49152
QY	1879	GTGGAGAAGTCCCTGAAGTAAGCAAGACHTTCTCTTAGTCGAGCCCAAGTTAAGAATG	1938
Db	49153	GTGGAGAAGTCCCTGAAGTAAGCAAGACHTTCTCTTAGTCGAGCCCAAGTTAAGAATG	49212
QY	1939	TTCTTATGTTGCCAGTGTCTTCTGTATGCTCATGCAAGCAGAGAACACATGGCGCTCTAGA	1998
Db	49213	TTCTTATGTTGCCAGTGTCTTCTGTATGCTCATGCAAGCAGAGAACACATGGCGCTCTAGA	49272
QY	1999	ACCAGGCAACTTTGGGAACTAGACTCCCAAGCTGGACTATGGCTCTACTTTTCAGGCCACAT	2058
Db	49273	ACCAGGCAACTTTGGGAACTAGACTCCCAAGCTGGACTATGGCTCTACTTTTCAGGCCACAT	49332
QY	2059	GGCTAAAGAAGTTTTCAGAAAGAAGTGGGACAGAGACAGAACTTTACCTTCATATATTTT	2118
Db	49333	GGCTAAAGAAGTTTTCAGAAAGAAGTGGGACAGAGACAGAACTTTACCTTCATATATTTT	49392
QY	2119	GTATGATCCTAATGAATGCATAAAATGTTTAAAGTTGATGGTGCATGAAATGTAATACTGTT	2178
Db	49393	GTATGATCCTAATGAATGCATAAAATGTTTAAAGTTGATGGTGCATGAAATGTAATACTGTT	49452
QY	2179	TTTAAACAACATGATTTGGAAATAAATCAATGCTATACATATGTTGATAAAG	2232
Db	49453	TTTAAACAACATGATTTGGAAATAAATCAATGCTATACATATGTTGATAAAG	49506

RESULT 7
 AAA35151
 ID AAA35151 standard; DNA; 149412 BP.
 XX
 AC AAA35151;
 XX
 DT 28-JUL-2000 (first entry)
 XX
 DE Human adenosine receptor related polynucleotide 2nd SEQ ID NO:25.
 XX
 KW Human; adenosine receptor; low adenosine antisense oligonucleotide;
 KW phosphorothioate; impaired respiration; inflammation; allergy;
 KW allergic disease; bronchoconstriction; inhibitor; antiinflammatory;
 KW antiallergic; antisthmatic; cytotatic; analgesic; impaired airway;
 KW lung disease; ischemic condition; pulmonary vasoconstriction; asthma;
 KW respiratory distress syndrome; pain; cystic fibrosis; emphysema;
 KW pulmonary hypertension; chronic obstructive pulmonary disease; COPD;
 KW cancer; leukaemia; lymphoma; carcinoma; metastasis; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200009525-A2.
 XX
 PD 24-FEB-2000.
 XX
 XX 03-AUG-1999; 99WO-US17712.
 PF
 XX 03-AUG-1998; 98US-0095212.
 PR
 XX (UYEC-) UNIV EAST CAROLINA.
 PA
 XX Nyce JW;
 PI
 XX WPI; 2000-205971/18.
 DR
 XX
 XX New antisense oligonucleotides useful for treating e.g. pulmonary
 PT vasoconstriction, inflammation, allergies, asthma, hypertension,
 PT bronchitis, emphysema, respiratory distress syndrome, ischemia or
 PT cancers -
 XX
 PS Disclosure; Page 1138-1171; 1343pp; English.
 XX
 CC The present invention describes a new composition comprising an
 CC antisense oligonucleotide (ON) with low adenosine (up to 15%), which
 CC targets nucleic acids involved in bronchoconstriction, allergies, and/or
 CC inflammation. The ON can have antiinflammatory, antiallergic,
 CC antisthmatic, cytotatic and analgesic activities. The compositions are
 CC useful for the treatment of diseases associated with inflammation,
 CC impaired airways, including lung disease and diseases whose secondary
 CC effects afflict the lungs of a subject. They can be used for treating
 CC e.g. ischaemic conditions, pulmonary vasoconstriction, allergies,
 CC asthma, impaired respiration, respiratory distress syndrome, pain, cystic
 CC fibrosis, pulmonary hypertension, emphysema, chronic obstructive
 CC pulmonary disease (COPD), and cancers such as leukaemias, lymphomas,
 CC carcinomas, and cancers which may metastasise to the lungs, including
 CC breast and prostate cancer. The reduction of the adenosine content of
 CC the ONs reduces side effects. The A-containing ONs break down with the
 CC release of deoxyadenosine which activates adenosine receptors causing
 CC bronchoconstriction and inflammation. AAA32313 to AAA3512 represent the
 CC nucleotide sequences given in the sequence listing from the present
 CC invention, which correspond to SEQ ID NO:1 to 2815, and then the last
 CC 195 sequences are also called SEQ ID NO:1 to 185, but the sequences
 CC differ from the previously named sequences. SEQ ID NO:11 to 1680
 CC (AAA32323 to AAA33992) are specifically claimed ONs from the present
 CC invention. N.B. Sequences given in the disclosure of the present
 CC invention do not match up with their corresponding SEQ ID NO: sequences
 CC given in the sequence listing.
 XX
 SQ Sequence 149412 BP; 43049 A; 31388 C; 33852 G; 41123 T; 0 other;

	Matches	1252;	Conservative	0;	Mismatches	2;	Indels	0;	Gaps	0;
Qy	979	AGCCTTTTTCACATAGCTCTTGGGTGTAGGATTGCCCACTCCAAAACCACTGTGTGA	1038							
Db	54597	AGCCTTTTTCACATAGCTCTTGGGTGTAGGATTGCCCACTCCAAAACCACTGTGTGA	54656							
Qy	1039	GGTCCAGAGTCAGACCAAGAAAGTGTGAAGTGACTACACAAGGACTCCTCGATGGT	1098							
Db	54657	GGTCCAGAGTCAGACCAAGAAAGTGTGAAGTGACTACACAAGGACTCCTCGATGGT	54716							
Qy	1099	CGTGAAGAAAGAACTCAATTGGCAGAGCCCTGAAGCCAGTCTTCAGACACAAGAGGA	1158							
Db	54717	CGTGAAGAAAGAACTCAATTGGCAGAGCCCTGAAGCCAGTCTTCAGACACAAGAGGA	54776							
Qy	1159	GCCTAGACACAAATGACAGATCTCTGCTTGGAAATCACACGCTCGCTTTCACAGATG	1218							
Db	54777	GCCTAGACACAAATGACAGATCTCTGCTTGGAAATCACACGCTCGCTTTCACAGATG	54836							
Qy	1219	TCTGATTACAGTGTGAATCTTGGTGTCTACGTTACCAGCAGGAGGCTGAGAGAGAG	1278							
Db	54837	TCTGATTACAGTGTGAATCTTGGTGTCTACGTTACCAGCAGGAGGCTGAGAGAGAG	54896							
Qy	1279	AGACTCCAGCTGGTTGGAAACAGTATTTCCAAACTACCTTCCAGTTCTCTCATTTTGG	1338							
Db	54897	AGACTCCAGCTGGTTGGAAACAGTATTTCCAAACTACCTTCCAGTTCTCTCATTTTGG	54956							
Qy	1339	AATACAGGCATAGATTACAGACTTTTAAATAGTAAATAATAATAATAATAATAATAA	1398							
Db	54957	AATACAGGCATAGATTACAGACTTTTAAATAGTAAATAATAATAATAATAATAATAA	55016							
Qy	1399	TGCAACTTGTAAATGTGGTAAAGAGTTAGTTGAGTTGCTATCATGTCAACCTGAAAT	1458							
Db	55017	TGCAACTTGTAAATGTGGTAAAGAGTTAGTTGAGTTGCTATCATGTCAACCTGAAAT	55076							
Qy	1459	GCTGTATTAGTCACAGAGATAATCTAGCTTTGAGCTTAAAGATTTTGAAGCAGTGGTAT	1518							
Db	55077	GCTGTATTAGTCACAGAGATAATCTAGCTTTGAGCTTAAAGATTTTGAAGCAGTGGTAT	55136							
Qy	1519	GTTTGGGAGACTGCTGAGTCAACCCCAATAGTTGTGATTTGGCAGAGGTTGGAAGTGTG	1578							
Db	55137	GTTTGGGAGACTGCTGAGTCAACCCCAATAGTTGTGATTTGGCAGAGGTTGGAAGTGTG	55196							
Qy	1579	ATCTGTGGGCACATTAGCCTATGTGCATGAGCATGAGTAAATGATGCTTTTGAATCA	1638							
Db	55197	ATCTGTGGGCACATTAGCCTATGTGCATGAGCATGAGTAAATGATGCTTTTGAATCA	55256							
Qy	1639	CAGTATACGCTCCATCCCTGCTCATCTCAGCTGGATCTCCATTCCTCAGGCTTGTGCGCA	1698							
Db	55257	CAGTATACGCTCCATCCCTGCTCATCTCAGCTGGATCTCCATTCCTCAGGCTTGTGCGCA	55316							
Qy	1699	AAAGCCCTTTTGTGTTTGTGTTTGTATGATGAGTATGAGTATGAGTATGAGTATGAGT	1758							
Db	55317	AAAGCCCTTTTGTGTTTGTGTTTGTATGATGAGTATGAGTATGAGTATGAGTATGAGT	55376							
Qy	1759	GTTCAGTGTCTCCAGATGCTTGTGATGCTCATATTTCTCCCTAAATTTGGCAGTGGAA	1818							
Db	55377	GTTCAGTGTCTCCAGATGCTTGTGATGCTCATATTTCTCCCTAAATTTGGCAGTGGAA	55436							
Qy	1819	CTCCTAAATCAAAATGGCTTCTAATCAAAGCTTTTAAACCCCTATTTGGTAAAGATGGAG	1878							
Db	55437	CTCCTAAATCAAAATGGCTTCTAATCAAAGCTTTTAAACCCCTATTTGGTAAAGATGGAG	55496							
Qy	1879	GTGAGAGAGCTCCCTGAGTAGCAAGACATTTTCCCTCTTAGTCGAGCCCAAGTTAAGAATG	1938							
Db	55497	GTGAGAGAGCTCCCTGAGTAGCAAGACATTTTCCCTCTTAGTCGAGCCCAAGTTAAGAATG	55556							
Qy	1939	TTCCTATGTCGCCAGTGTGTTTCTGATCTGATGCAAGCAAGAAACACTGGGCTTCTAGA	1998							
Db	55557	TTCCTATGTCGCCAGTGTGTTTCTGATCTGATGCAAGCAAGAAACACTGGGCTTCTAGA	55616							
Qy	1999	ACCAGGCAACTGGGAACCTAGACTCCCAAGCTGGACTATGGCTCTACTTTTCAGGCCACAT	2058							
Db	55617	ACCAGGCAACTGGGAACCTAGACTCCCAAGCTGGACTATGGCTCTACTTTTCAGGCCACAT	55676							

Query Match 51.6%; Score 1152; DB 21; Length 149412;
 Best Local Similarity 99.8%; Pred. No. 0;

QY 2059 GGCTAAAGAGGTTTCAGAAAGAGTGGGACAGACAGAACTTTCACCTTCATATATTT 2118
|||||
Db 55677 GGCTAAAGAGGTTTCAGAAAGAGTGGGACAGACAGAACTTTCACCTTCATATATTT 55736
QY 2119 GTATGATCCCTAATGAATGCATATAAATGTAAGTTGATGGTGAATGTAATGTAATGTT 2178
|||||
Db 55737 GTATGATCCCTAATGAATGCATATAAATGTAAGTTGATGGTGAATGTAATGTAATGTT 55796
QY 2179 TTTAACTAATGATTTGGAAATATAATCAATGCTATATACTATGTTGATAAAG 2232
|||||
Db 55797 TTTAACTAATGATTTGGAAATATAATCAATGCTATATACTATGTTGATAAAG 55850

RESULT 8

AAF21273

ID AAF21273 standard; DNA; 152740 BP.

AC AAF21273;

XX 14-MAR-2001 (first entry)

DT 14-MAR-2001 (first entry)

XX Human low adenosine antisense oligonucleotide related sequence #2840.

DE Low adenosine antisense oligonucleotide; phosphorothioate; allergy;
KW human; airway disorder; bronchoconstriction; lung inflammation;
KW surfactant depletion; respiratory; bronchodilator; antiinflammatory;
KW immunosuppressive; antiasthmatic; analgesic; hypotensive; cytostatic;
KW respiratory obstruction; pulmonary obstruction; impeded respiration;
KW surfactant hypoproduction; pulmonary vasoconstriction; asthma; RDS;
KW respiratory distress syndrome; pulmonary pain; cystic fibrosis; allergic rhinitis;
KW pulmonary hypertension; emphysema; pulmonary transplantation rejection;
KW chronic obstructive pulmonary disease; pulmonary infection; bronchitis;
KW cancer; ss.

XX Homo sapiens.

OS Homo sapiens.

FN WO200062736-A2.

PN 26-OCT-2000.

PD 26-OCT-2000.

PP 24-MAR-2000; 2000WO-US08020.

PF 06-APR-1999; 99US-0127958.

PR (UYEC-) UNIV EAST CAROLINA.

XX (NYCE/) NYCE J W.

PA NYCE J W.

PI Nyce JW;

XX WPI; 2000-679539/66.

DR WPI; 2000-679539/66.

XX Low adenosine (A) content antisense oligonucleotides which do not

XX trigger adenosine receptors during metabolism, useful e.g. for treating

XX cancers and respiratory obstructions.

PT Disclosure; Page 1219-1254; 1592pp; English.

XX The present invention describes low adenosine (A) content antisense

XX oligonucleotides and compositions (I) comprising them. In the antisense

XX oligonucleotides the A is replaced by a 'Universal' or alternative base.

XX (I) can have respiratory, bronchodilator, antiinflammatory, analgesic,

XX immunosuppressive, antiasthmatic, hypotensive and cytostatic activities.

XX The antisense oligonucleotides and (I) can be used to down-regulate the

XX expression and/or activity of target polypeptides associated with

XX lung/respiratory disorders and malignancies, such as stimulating and

XX activating peptide factors and transmitters, antibody receptors, cytokines and

XX immunoglobulins and antibodies, transcription factors,

XX chemokines, endogenously produced specific and non-specific enzymes,

XX binding proteins, adhesion molecules and their receptors, cytokine and

XX chemokine receptors, adenosine receptors, bradykinin receptors, central

XX nervous system (CNS) and peripheral nervous and non-nervous system

XX receptors, CNS and peripheral nervous and non-nervous system peptide

XX

XX

XX

XX

XX

XX

XX

XX

transmitters, defensins, growth factors, vasoactive peptides and
receptors, binding proteins and malignancy associated proteins. The
antisense oligonucleotides may be used in this way to treat disorders
including respiratory obstruction (especially pulmonary obstruction
and/or bronchoconstriction) and/or lung inflammation, allergy(ies)
and/or surfactant hypoproduction which are associated with a disease or
condition selected from pulmonary vasoconstriction, inflammation,
allergies, asthma, impeded respiration, respiratory distress syndrome
(RDS), pain, cystic fibrosis (CF), allergic rhinitis (AR), pulmonary
hypertension, emphysema, chronic obstructive pulmonary disease (COPD),
pulmonary transplantation rejection, pulmonary infections, bronchitis,
and/or cancer. AAF18434 to AAF21543 represent human polynucleotide
fragments and antisense oligonucleotides used in the exemplification of
the present invention.

XX

SQ Sequence 152740 BP; 44169 A; 32023 C; 34549 G; 41999 T; 0 other;

Query Match 51.6%; Score 1152; DB 21; Length 152740;

Best Local Similarity 99.8%; Pred. No. 0;

Matches 1252; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 979 AGCCTTTTTCACATAGCTCTTGGCTGTAGGATTGCCCTCCCAAAACACAGTGTGTGGA 1038

Db 54597 AGCCTTTTTCACATAGCTCTTGGCTGTAGGATTGCCCTCCCAAAACACAGTGTGTGGA 54656

QY 1039 GGTCCAGGAGTGAGACAGGAGAAAGATGTAAAGTGACTACACAGGACTCTCGATGGT 1098

Db 54657 GGTCCAGGAGTGAGACAGGAGAAAGATGTAAAGTGACTACACAGGACTCTCGATGGT 54716

QY 1099 CGTGGAAAGGAAAGTCAATTCGACAGAGCCCTGAGCCAGTCTTCAGGACAAGAGGA 1158

Db 54717 CGTGGAAAGGAAAGTCAATTCGACAGAGCCCTGAGCCAGTCTTCAGGACAAGAGGA 54776

QY 1159 GCCTAGACAGAAATGACAGATCTCTGCTTTGGAATCACACAGCTGCTGCTTCCACAGAT 1218

Db 54777 GCCTAGACAGAAATGACAGATCTCTGCTTTGGAATCACACAGCTGCTGCTTCCACAGAT 54836

QY 1219 TGTGATTCACAGTGTGAATCTTGGTGTCTACGTTACCAGGAGGAGGAGAGAG 1278

Db 54837 TGTGATTCACAGTGTGAATCTTGGTGTCTACGTTACCAGGAGGAGGAGAGAG 54896

QY 1279 AGACTCCAGCTGGGTGGGAAACAGTATTTCCAAACTACCTTCCAGTTCCTCATTTTG 1338

Db 54897 AGACTCCAGCTGGGTGGGAAACAGTATTTCCAAACTACCTTCCAGTTCCTCATTTTG 54956

QY 1339 AATACAGGCATAGAGTTCAGACTTTTAAATAGTAAATAAATAAGCTGAAAC 1398

Db 54957 AATACAGGCATAGAGTTCAGACTTTTAAATAGTAAATAAATAAGCTGAAAC 55016

QY 1399 TGCAACTTGTAAATGTGTAAAGAGTGTAGTTGCTTCTCATGTCACAAAGCTGAAAT 1458

Db 55017 TGCAACTTGTAAATGTGTAAAGAGTGTAGTTGCTTCTCATGTCACAAAGCTGAAAT 55076

QY 1459 GCTGTATTAGTCACAGAGATAATTTCTAGCTTTGAGCTTAAAGAAATTTTTCAGAGGTTGGTAT 1518

Db 55077 GCTGTATTAGTCACAGAGATAATTTCTAGCTTTGAGCTTAAAGAAATTTTTCAGAGGTTGGTAT 55136

QY 1519 GTTTGGAGAGTCTGCTAGTCAACCAATAGTTGTTGATTGGCAGAGTGTGAAAGTGTGTG 1578

Db 55137 GTTTGGAGAGTCTGCTAGTCAACCAATAGTTGTTGATTGGCAGAGTGTGAAAGTGTGTG 55196

QY 1579 ATCTGGGACATTAAGCTTATGTCATGACGATCTAAAGTAATGATGTCCTTTGAATCA 1638

Db 55197 ATCTGGGACATTAAGCTTATGTCATGACGATCTAAAGTAATGATGTCCTTTGAATCA 55256

QY 1639 CAGTATACCTCCATCGCTCATCTCAGCTGGATCTCCATCTCTCAGGCTTGTGCTGCA 1698

Db 55257 CAGTATACCTCCATCGCTCATCTCAGCTGGATCTCCATCTCTCAGGCTTGTGCTGCA 55316

QY 1699 AAAGCCTTTTGTGTTTTTGTATCATATTATGAAGTCATCGCTTAAATACATTCAGT 1758

Db 55317 AAAGCCTTTTGTGTTTTTGTATCATATTATGAAGTCATCGCTTAAATACATTCAGT 55376

Mon Jun 2 09:42:04 2003

1759 GTTTCAGTGTTCGAGATGCTCCTGATGCTCATATTTGTTCCCTAATTTCCAGTGGAA 1818
 55377 GTTTCAGTGTTCGAGATGCTCCTGATGCTCATATTTGTTCCCTAATTTCCAGTGGAA 55436
 1819 CTCTAATCAAAATGGCTTCTAATCAAAAGCTTTTAAACCTATTTGGTAAAGATGGAAG 1878
 55437 CTCTAATCAAAATGGCTTCTAATCAAAAGCTTTTAAACCTATTTGGTAAAGATGGAAG 55496
 1879 GTGAGAGAGCTCCCTGAAGTAAGCAAAAGACTTTCTCTTAGTCGAGCAAGTTAAGATG 1938
 55497 GTGAGAGAGCTCCCTGAAGTAAGCAAAAGACTTTCTCTTAGTCGAGCAAGTTAAGATG 55556
 1939 TTCTTATGTTCCAGAGTGTCTTCTGATCTGATCAAGCAAGCAACACTTGGGCTTCTAGA 1998
 55557 TTCTTATGTTCCAGAGTGTCTTCTGATCTGATCAAGCAAGCAACACTTGGGCTTCTAGA 55616
 1999 ACCAGGCAACTTGGGAAGTACAGTCCCAAGCTGACATGCTTACCTTTTACAGCCACAT 2058
 55617 ACCAGGCAACTTGGGAAGTACAGTCCCAAGCTGACATGCTTACCTTTTACAGCCACAT 55676
 2059 GGCTAAAGAGAGTTTCAGAAAGAGTGGGGACAGAGCAAGACTTTTACCTTCTATATTT 2118
 55677 GGCTAAAGAGAGTTTCAGAAAGAGTGGGGACAGAGCAAGACTTTTACCTTCTATATTT 55736
 2119 GTATGATCCTAATGAATGATCAAAATGTTAAGTTGATGATGATGATGATGATGATGAT 2178
 55737 GTATGATCCTAATGAATGATCAAAATGTTAAGTTGATGATGATGATGATGATGATGAT 55796
 2179 TTTTAACTATGATTTGGAAATTAATCAATGCTATATATGTTGATAAAAG 2232
 55797 TTTTAACTATGATTTGGAAATTAATCAATGCTATATATGTTGATAAAAG 55850

RESULT 9
 AAQ96298 standard; cDNA; 1979 BP.

AAQ96298;
 29-DEC-1995 (first entry)
 Human monocyte chemoattractant protein-1 receptor MCP-1RB.
 Monocyte chemoattractant protein-1 receptor; MCR-1R; chemokine; ss.
 Homo sapiens.
 Key Location/Qualifiers
 CDS 81..1160
 /*tag= a
 W09519436-A.
 20-JUL-1995.
 11-JAN-1995; 95WO-US00476.
 13-JAN-1994; 94US-0182962.
 (REGC) UNIV CALIFORNIA.
 Charo I, Coughlin S;
 WPI; 1995-263866/34.
 P-PSDB; AAR79166.
 DNA encoding monocyte chemo-attractant protein-1 receptor - used partic.
 for identifying antagonists and for treating diseases characterised by
 monocytic infiltrates
 Disclosure; Fig 2; 84pp; English.
 To identify and clone new members of the chemokine receptor gene

family, degenerate oligo primers were designed corresp. to the conserved sequences R79167 in the second and R79168 in the third transmembrane domains of the MIP-lalpha/RANTES receptor, the IL-8 receptors and the HUMSTRS orphan receptor (Genbank Accession #M99293). The degenerate oligo incorporating EcoRI and XhoI sites at their 5' ends are Q96299 and Q96300. Amplification of cDNA derived from MM6 cells with the primers yielded a number of PCR products. One cDNA appeared to encode a novel protein. To obtain a full-length version of this clone, a MM6 cDNA library was constructed in pFROG and probed with the PCR product. A 2.1 kb cDNA library revealed a second additional clones in the MM6 cDNA library revealed a second sequence that was identical to the 2.1 kb cDNA sequence first obtd. from the 5' UTR through the putative seventh transmembrane domain but contained a different cytoplasmic tail. The second sequence appears to represent alternative splicing of the carboxyl-terminal tail of the MCP-1R protein. The two sequences are denoted MCP-1RA and MCP-1RB (see Q96297/R79165 & Q96298/R79166). Active mature MCP-1RA has a predicted mol. wt. of about 42,000 daltons. MCP-1RB has a mol. wt. of about 41,000 daltons.

Sequence 1979 BP; 530 A; 434 C; 452 G; 563 T; 0 other;

Query Match	43.9%;	Score 980;	DB 16;	Length 1979;
Best Local Similarity	100.0%;	Pred. No. 0;		
Matches 980;	Conservative	0;	Mismatches	0;
Indels	0;	Gaps	0;	

QY 1 GGATTGAACAAGACGATTTCCCAAGTACATCCCAACATGCTGTCCACATCTCGTTCT 60
 DB 42 GGATTGAACAAGACGATTTCCCAAGTACATCCCAACATGCTGTCCACATCTCGTTCT 101
 QY 61 CGGTTTATCAGAAATACCAACGAGCGGTGAAGAAGTACCAACCTTTTGTGATTATGAT 120
 DB 102 CGGTTTATCAGAAATACCAACGAGCGGTGAAGAAGTACCAACCTTTTGTGATTATGAT 161
 QY 121 TAGCGTCTCCCTGTCATAAATTTGACGTGAAGCAAAATTTGGGCCCAACTCTCGCTCCG 180
 DB 162 TAGCGTCTCCCTGTCATAAATTTGACGTGAAGCAAAATTTGGGCCCAACTCTCGCTCCG 221
 QY 181 CTCTACTCGCTGGTGTTCATCTTTGGTTTGGGCAACATGCTGGTCTCCATCTTA 240
 DB 222 CTCTACTCGCTGGTGTTCATCTTTGGTTTGGGCAACATGCTGGTCTCCATCTTA 281
 QY 241 ATAACTGCAAAAGCTGAAGTCTTACCTGACATGACATTTACCTGCTCAACCTGCCATCTCT 300
 DB 282 ATAACTGCAAAAGCTGAAGTCTTACCTGACATGACATTTACCTGCTCAACCTGCCATCTCT 341
 QY 301 GATCTGCTTTTCTTATTACTCTCCCATTTGGGCTCACTCTGCTGCAAAATGAGTGGGTC 360
 DB 342 GATCTGCTTTTCTTATTACTCTCCCATTTGGGCTCACTCTGCTGCAAAATGAGTGGGTC 401
 QY 361 TTTGGGAATGCAATGTGCAAAATTTTACAGGGCTGTATCATCGGTTATTTTGGCGGA 420
 DB 402 TTTGGGAATGCAATGTGCAAAATTTTACAGGGCTGTATCATCGGTTATTTTGGCGGA 461
 QY 421 ATCTCTTCATCATCTCTCTGACATGATAGATACCTGGCTATTGTCCATGCTGTGTTT 480
 DB 462 ATCTCTTCATCATCTCTCTGACATGATAGATACCTGGCTATTGTCCATGCTGTGTTT 521
 QY 481 GCTTTAAAGCCAGGACGGTCACTTTGGGGTGGTGACAAGTGTATCATCACCTGGTGGTG 540
 DB 522 GCTTTAAAGCCAGGACGGTCACTTTGGGGTGGTGACAAGTGTATCATCACCTGGTGGTG 581
 QY 541 GCTGTGTTTGTCTTCTCCAGGAATCATCTTTTACTAAATGCCAGAAAGATTTCTGTT 600
 DB 582 GCTGTGTTTGTCTTCTCCAGGAATCATCTTTTACTAAATGCCAGAAAGATTTCTGTT 641
 QY 601 TATGTCTGTGGCCCTTATTTTCCAGGAGATGGAATAATTTCCACACATATAGGAGAC 660
 DB 642 TATGTCTGTGGCCCTTATTTTCCAGGAGATGGAATAATTTCCACACATATAGGAGAC 701
 QY 661 ATTTTGGGCTGGTCTGCCGTGCTCATCATGCTGCTACTCGGGAATCCCTGAA 720
 DB 702 ATTTTGGGCTGGTCTGCCGTGCTCATCATGCTGCTACTCGGGAATCCCTGAA 761

ID AAS12139 standard; DNA: 1083 BP.
 AC AAS12139;
 XX 04-DEC-2001 (first entry)
 DT Human CCR2-641 polymorphic variant polynucleotide.
 XX Human; CCR2 receptor; CCR2-64V; gene therapy; atherosclerosis;
 XX single nucleotide polymorphism; hypercholesterolaemia; ds.
 KW Homo sapiens.
 OS
 XX Key Location/Qualifiers
 XX variation replace(190,G)
 FT /*tag= a
 FT /standard_name= "Single nucleotide polymorphism"
 XX WO200162796-A1.
 PN 30-AUG-2001.
 PD 22-FEB-2001; 2001WO-GB00755.
 XX 22-FEB-2000; 2000GB-0004183.
 PR (SMIK) SMITHKLINE BEECHAM PLC.
 PA Valdes AM, Groot PHE, Spurr NK;
 PI WPI; 2001-550086/61.
 XX P-PSDB; AAU07613.
 DR Diagnosing atherosclerosis or susceptibility to atherosclerosis in a
 XX subject, by determining a single nucleotide polymorphism in specific
 PT codon of a polynucleotide encoding human CCR2 receptor in genome of the
 PT subject -
 XX Claim 3; Page 20; 28pp; English.
 PS The invention relates to diagnosing atherosclerosis (or susceptibility
 XX to) in a subject by determining expression or activity of the human
 CC CCR2-641 polypeptide (a polymorphic variant form of the human CCR2
 CC receptor) or the CCR2-64V polypeptide (human CCR2 receptor), by screening
 CC for a single nucleotide polymorphism in codon 64 of the polynucleotide
 CC encoding the CCR2 receptor. This results in production of CCR2-64L,
 CC whereby polymorphic variants are associated with a lower incidence of
 CC atherosclerosis. The presence or amount of CCR2-64I/V in a sample can
 CC also be analysed. The sequences of the invention can be used for
 CC predicting the response of a patient to drug treatment, for predicting
 CC the disease outcome in a patient and also for the production of a
 CC treatment for hypercholesterolaemia. The sequence represents DNA encoding
 CC the polymorphic variant polypeptide CCR2-64I.
 XX Sequence 1083 BP; 256 A; 260 C; 246 G; 321 T; 0 other;
 SQ
 Query Match 39.98; Score 890; DB 22; Length 1083;
 Best Local Similarity 99.98; Pred. No. 0;
 Matches 940; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 40 ATGCTGTCACATCTCGTTTCGGTTTATCAGAAATACCAACGAGAGCGGTGAAGATC 99
 DB 1 ATGCTGTCACATCTCGTTTCGGTTTATCAGAAATACCAACGAGAGCGGTGAAGATC 60
 QY 100 ACCACCTTTTGGATTATGATTACGGTCTCCCTCTCATAAATTTGACGTGAAGCAAT 159
 DB 61 ACCACCTTTTGGATTATGATTACGGTCTCCCTCTCATAAATTTGACGTGAAGCAAT 120
 QY 160 GGGGCCCAACTCCTCGCTCCGCTACTCGCTGGTGTTCATCTTTGGTTTGTGGCAAC 219
 DB 121 GGGGCCCAACTCCTCGCTCCGCTACTCGCTGGTGTTCATCTTTGGTTTGTGGCAAC 180
 QY 220 ATGCTGGTCTCTCATCTTAATAAAGCTGCAAAAGCTGAAGTGTGACTGACATTTAC 279

DB 181 ATGCTGGTCATCCTCATCTTAATAAAGCTGAAAAGCTGAAGTCTTGACTGACATTTAC 240
 QY 280 CTGCTCAACCTGGCCATCTCTGATCTGCTTTTCTTATCTACTCTCCCATTTGGGCTCAC 339
 DB 241 CTGCTCAACCTGGCCATCTCTGATCTGCTTTTCTTATCTACTCTCCCATTTGGGCTCAC 300
 QY 340 TCTGCTGCAAAATGAGTGGTCTTTTGGGAATGCAATGTGCAAAATTTATTCACAGGGTGTAT 399
 DB 301 TCTGCTGCAAAATGAGTGGTCTTTTGGGAATGCAATGTGCAAAATTTATTCACAGGGTGTAT 360
 QY 400 CACATCGGTTATTTTGGGGAATCTTCTTCATCATCTCTCTGACATTCGATGATACCTG 459
 DB 361 CACATCGGTTATTTTGGGGAATCTTCTTCATCATCTCTCTGACATTCGATGATACCTG 420
 QY 460 GCTATTTGCCATGCTGCTTTTAAAGGACGAGCGGTACCTTTGGGGTGGTGACA 519
 DB 421 GCTATTTGCCATGCTGCTTTTAAAGGACGAGCGGTACCTTTGGGGTGGTGACA 480
 QY 520 AGTGTGATCACCTGGTGGTGGTGTGTTTGTGTTCTCCAGGAATCATCTTTACTATA 579
 DB 481 AGTGTGATCACCTGGTGGTGGTGTGTTTGTGTTCTCCAGGAATCATCTTTACTATA 540
 QY 580 TGCCAGAAAGAGATTCTGTTTATGCTGTGGCCCTTATTTCCACGAGATGGAATAAT 639
 DB 541 TGCCAGAAAGAGATTCTGTTTATGCTGTGGCCCTTATTTCCACGAGATGGAATAAT 600
 QY 640 TTCCACACAATAATGAGGAACATTTTGGGCTGTCTCTCGCTGTCTCATCTGTCATC 699
 DB 601 TTCCACACAATAATGAGGAACATTTTGGGCTGTCTCTCGCTGTCTCATCTGTCATC 660
 QY 700 TGCTACTCGGGAATCTCTGAAACCCCTGCTGGTGTGTCGAAACGAGAGAGGAGCATAGG 759
 DB 661 TGCTACTCGGGAATCTCTGAAACCCCTGCTGGTGTGTCGAAACGAGAGAGGAGCATAGG 720
 QY 760 GCAGTGAGATCATCTTCACATCATGATTTTACTTTCTTCTTCTGACTCCCTATAAC 819
 DB 721 GCAGTGAGATCATCTTCACATCATGATTTTACTTTCTTCTTCTGACTCCCTATAAC 780
 QY 820 ATTGTCAATCTCTCTGAAACCCCTGCTGGTGTGTCGAAACGAGAGAGGAGCATAGG 879
 DB 781 ATTGTCAATCTCTCTGAAACCCCTGCTGGTGTGTCGAAACGAGAGAGGAGCATAGG 840
 QY 880 AGTCAACTGGACCAAGCCACGAGTGCACAGAGACTCTTGGGATGACTCACTGCTGTCATC 939
 DB 841 AGTCAACTGGACCAAGCCACGAGTGCACAGAGACTCTTGGGATGACTCACTGCTGTCATC 900
 QY 940 AATCCCATCATCTATGCTTCTGTTGGGGAAGTTTCAGAAG 980
 DB 901 AATCCCATCATCTATGCTTCTGTTGGGGAAGTTTCAGAAG 941
 RESULT 12
 AAT96976
 ID AAT96976 standard; cDNA: 1083 BP.
 XX AAT96976;
 AC AAT96976;
 XX 27-FEB-1998 (first entry)
 DT Human monocyte chemoattractant protein 1 receptor encoding cDNA.
 XX Human; MCP-1; monocyte chemoattractant protein; receptor; tumour;
 KW inflammatory disease; viral; allergy; diabetes; ds.
 OS Homo sapiens.
 XX Key Location/Qualifiers
 XX CDS 1..1083
 FT /*tag= a
 FT /product= Monocyte_chemoattractant_protein_1_receptor
 XX JP09238688-A.

XX PD 16-SEP-1997.
 XX PF 11-MAR-1996; 96JP-0053574.
 XX PR 11-MAR-1996; 96JP-0053574.
 XX PA (TAKE) TAKEDA CHEM IND LTD.
 DR WPI; 1997-506557/47.
 DR P-PSDB; RAW35833.
 XX
 PT DNA encoding human monocyte chemoattractant protein 1 receptor -
 PT used to treat tumours and inflammatory, viral, infectious, allergic,
 PT diabetic and central nervous system diseases
 XX
 PS Claim 1; Page 12; 15pp; Japanese.
 XX
 CC The present sequence encodes human monocyte chemoattractant protein 1
 CC (MCP-1) receptor protein. The MCP-1 receptor protein and encoding DNA
 CC are used for the prevention and treatment of tumours and inflammatory,
 CC viral, infectious, allergic, diabetic and central nervous system
 CC diseases.
 XX
 SQ Sequence 1083 BP; 257 A; 259 C; 245 G; 322 T; 0 other;
 Query Match 37.6%; Score 839; DB 18; Length 1083;
 Best Local Similarity 99.8%; Pred. No. 0;
 Matches 939; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

 QY 40 ATGCTGTCCACATCTCGTCTCGTGTATCAGAAATACCAACGAGCGGTGAAGAGTC 99
 DB 1 ATGCTGTCCACATCTCGTCTCGTGTATCAGAAATACCAACGAGCGGTGAAGAGTC 60
 QY 100 ACCACCTTTTTCATATGATGATGATGATGATGATGATGATGATGATGATGAT 159
 DB 61 ACCACCTTTTTCATATGATGATGATGATGATGATGATGATGATGATGATGAT 120
 QY 160 GGGGCCCAACTCTGCTCGCTCTACTCGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 219
 DB 121 GGGGCCCAACTCTGCTCGCTCTACTCGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 180
 QY 220 ATGCTGTGCTGCTCTCATCTTAATACTGCAAAAGCTGAACTGCTGCTGCTGCTGCT 279
 DB 181 ATGCTGTGCTGCTCTCATCTTAATACTGCAAAAGCTGAACTGCTGCTGCTGCTGCT 240
 QY 280 CTGCTCAACCTGGCCATCTCTGATCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 339
 DB 241 CTGCTCAACCTGGCCATCTCTGATCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 300
 QY 340 TCTGCTGCAATGATGCTGGTCTTTGGGAATGCAATGCTGCAATGCTGCAATGCTGCAAT 399
 DB 301 TCTGCTGCAATGATGCTGGTCTTTGGGAATGCAATGCTGCAATGCTGCAATGCTGCAAT 360
 QY 400 CACATCGGTTATTTGGCGGAATCTTCTTATCATCTCTGCAATGCTGCAATGCTGCAAT 459
 DB 361 CACATCGGTTATTTGGCGGAATCTTCTTATCATCTCTGCAATGCTGCAATGCTGCAAT 420
 QY 460 GCTATGTCATGCTGCTGCTTTTAAAGCCAGGAGGCTGCTGCTGCTGCTGCTGCTGCT 519
 DB 421 GCTATGTCATGCTGCTGCTTTTAAAGCCAGGAGGCTGCTGCTGCTGCTGCTGCTGCT 480
 QY 520 AGTGTGATCACTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 579
 DB 481 AGTGTGATCACTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 540
 QY 580 TGGCAGAAAGAGATCTGTTTATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 639
 DB 541 TGGCAGAAAGAGATCTGTTTATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 600
 QY 640 TTCCACACATATAGGAGCAATTTTGGGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 699
 DB 601 TTCCACACATATAGGAGCAATTTTGGGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 660

QY 700 TGCTACTCGGAATCCTGAAACCCCTGCTTGGTGTGCAACGAGAGAGGCGATAGG 759
 DB 661 TGCTACTCGGAATCCTGAAACCCCTGCTTGGTGTGCAACGAGAGAGGCGATAGG 720
 QY 760 GCAGTGAGAGTCACTTCCACCATCATGATGTTTACTTCTTCTTGGACTCCCTATAAT 819
 DB 721 GCAGTGAGAGTCACTTCCACCATCATGATGTTTACTTCTTCTTGGACTCCCTATAAT 780
 QY 820 ATTGTCAATTCCTGCAACACCTTCCAGGAATTCCTTGGCTGAGTAACCTGCTGATC 879
 DB 781 ATTGTCAATTCCTGCAACACCTTCCAGGAATTCCTTGGCTGAGTAACCTGCTGATC 840
 QY 880 AGTCAACTGGACCAAGCCAGCGACAGAGACTCTTGGGATGACTCACTGCTGCTGATC 939
 DB 841 AGTCAACTGGACCAAGCCAGCGACAGAGACTCTTGGGATGACTCACTGCTGCTGATC 900
 QY 940 AATCCCATCATCTATGCTTGGTGGGAGAGTTCAGAAG 980
 DB 901 AATCCCATCATCTATGCTTGGTGGGAGAGTTCAGAAG 941

RESULT 13

ABI97976
 ID ABI97976 standard; cDNA; 1083 BP.
 AC ABI97976;
 XX
 DT 18-FEB-2002 (first entry)
 XX
 DE Non-endogenous human GPCR cDNA, SEQ ID NO: 472.
 XX
 KW Human; G protein-coupled receptor; GPCR; non-endogenous; mutant;
 XX constitutively activated GPCR; agonist; disease; ss.
 OS Homo sapiens.
 OS Synthetic.
 XX
 PN WO200177172-A2.
 XX
 PD 18-OCT-2001.
 XX
 PF 05-APR-2001; 2001WO-US11098.
 XX
 PR 07-APR-2000; 2000US-195747P.
 XX
 PA (AREN-) ARENA PHARM INC.
 XX
 PI Lehmann-Bruinsma K, Liaw CW, Lin I;
 XX
 DR WPI; 2001-648759/74.
 DR P-PSDB; ABB56340.
 XX

Identifying agonists of G protein-coupled receptors (GPCRs) for use in
 disease treatment, comprises contacting candidate compounds with
 versions of GPCRs -

Example 2; Page 273-274; 394pp; English.

The invention relates to G protein-coupled receptors (GPCRs) for which
 the endogenous ligand has been identified. Non-endogenous
 constitutively activated versions of known GPCRs are used in the
 invention for the direct identification of candidate compounds as
 receptor agonists, inverse agonists or partial agonists. Such
 agonists are useful as therapeutic agents for diseases or disorders
 associated with GPCRs. The present sequence encodes a non-endogenous
 version of a known human GPCR.

Sequence 1083 BP; 257 A; 260 C; 246 G; 320 T; 0 other;

Query Match

Best Local Similarity 99.8%; Score 839; DB 23; Length 1083;

Matches 939; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY	40	ATGCTGCCACATCTCGTTCTCGGTTTATCAGAAATACCAACGAGAGCGGTGAAGAATC	99
DB	1	ATGCTGCCACATCTCGTTCTCGGTTTATCAGAAATACCAACGAGAGCGGTGAAGAATC	60
QY	100	ACCACCTTTTGTGATTATGATTACGGTGCTCCCTGTCTATAAATTTGACGTGAAGCAATTT	159
DB	61	ACCACCTTTTGTGATTATGATTACGGTGCTCCCTGTCTATAAATTTGACGTGAAGCAATTT	120
QY	160	GGGGCCCAACTCCGCTCCGCTCTACATCGTGGTGTTCATCTTTGGTTTCTTGGGCAAC	219
DB	121	GGGGCCCAACTCCGCTCCGCTCTACATCGTGGTGTTCATCTTTGGTTTCTTGGGCAAC	180
QY	220	ATGCTGTGTCGCTCATCTTATAAAGCTGCAAAAGCTGAACTGCTTGACTGACATTTAC	279
DB	181	ATGCTGTGTCGCTCATCTTATAAAGCTGCAAAAGCTGAACTGCTTGACTGACATTTAC	240
QY	280	CTGCTCAACCTGGCCATCTCTGATCTGCTTTTCTTATTTACATCTCCCATTTGGGCTCAC	339
DB	241	CTGCTCAACCTGGCCATCTCTGATCTGCTTTTCTTATTTACATCTCCCATTTGGGCTCAC	300
QY	340	TCCTGCTGCAAAATGAGTGGGTCTTTGGGAATGCAATGTCAAAATTAATTCAGGGCTGTAT	399
DB	301	TCCTGCTGCAAAATGAGTGGGTCTTTGGGAATGCAATGTCAAAATTAATTCAGGGCTGTAT	360
QY	400	CACATCGGTTATTTGGCGGAATCTTCTTCATCATCTCTCTGACATCGATAGATACCTG	459
DB	361	CACATCGGTTATTTGGCGGAATCTTCTTCATCATCTCTCTGACATCGATAGATACCTG	420
QY	460	GCTATTGTCATCTGCTGTTTGCTTTAAAAGCCAGACGGTCACTTTGGGTTGGTGACA	519
DB	421	GCTATTGTCATCTGCTGTTTGCTTTAAAAGCCAGACGGTCACTTTGGGTTGGTGACA	480
QY	520	AGTGTGATCACCTGGTGGTGGTGTGTTTGCTCTCTGCCAGGAATCATCTTTACTAAA	579
DB	481	AGTGTGATCACCTGGTGGTGGTGTGTTTGCTCTCTGCCAGGAATCATCTTTACTAAA	540
QY	580	TGCCAGAAAGAAATCTGTTTATGTCTGTGGCCCTTATTTTCCAGAGGATGGAATAAT	639
DB	541	TGCCAGAAAGAAATCTGTTTATGTCTGTGGCCCTTATTTTCCAGAGGATGGAATAAT	600
QY	640	TTCCACACAATAATGAGGAACATTTTGGGCTGGTCTCGCGTGCCTCATCATGGTCAATC	699
DB	601	TTCCACACAATAATGAGGAACATTTTGGGCTGGTCTCGCGTGCCTCATCATGGTCAATC	660
QY	700	TGCTACTCGGGAATCCGTGAAACCCCTGCTCGGTGTGAAACGAGAGAAGAGGCATAGG	759
DB	661	TGCTACTCGGGAATCCGTGAAACCCCTGCTCGGTGTGAAACGAGAGAAGAGGCATAGG	720
QY	760	GCACTGAGAGTCATCTTCCACCATCATGATGTTTACTTCTCTCTGACCTCCCTATTAAC	819
DB	721	GCAAGAGAGTCATCTTCCACCATCATGATGTTTACTTCTCTCTGACCTCCCTATTAAC	780
QY	820	ATTCTCATCTCCTGAACACCTTCCAGGAATTTCTCGGCCTGAGTAACCTGTGAAAGCAC	879
DB	781	ATTGTTCATCTCCTGAAACACCTTCCAGGAATTTCTTCCGCCCTGAGTAACCTGTGAAAGCAC	840
QY	880	AGTCAACTGGACCAAGCCAGCGAGTGCAGAGACTCTTGGGATGACTCAGTCTGCTGCATC	939
DB	841	AGTCAACTGGACCAAGCCAGCGAGTGCAGAGACTCTTGGGATGACTCAGTCTGCTGCATC	900
QY	940	AATCCCATCATCTATGCCCTTCGTTGGGGAAGAGTTTCAGAAAG	980
DB	901	AATCCCATCATCTATGCCCTTCGTTGGGGAAGAGTTTCAGAAAG	941
RESULT	14		
AAV84136			
ID	AAV84136	standard; DNA; 461 BP.	
XX			
AC	AAV84136;		
XX			
DT	15-MAR-1999	(first entry)	

XX	HIV-1 co-receptor CCR5 gene hybridisation probe.
DE	
XX	HIV-1; CCR5; CCR5m303; co-receptor; infection; diagnosis; AIDS;
KW	gene therapy; human; probe; ss.
OS	Synthetic.
OS	Homo sapiens.
OS	
XX	WO9854317-A1.
PN	
XX	03-DEC-1998.
PD	
XX	29-MAY-1998; 98WO-EP03437.
PF	
XX	30-MAY-1997; 97US-0048057.
PR	
XX	(MOND-) FOND MONDIALE RECH & PREVENTION SIDA.
PA	
XX	Arenzana Slesdedos F, Beretta A, Braun J, Quillent C;
PI	
XX	WPI; 1999-059835/05.
DR	
XX	New CCR5 variant protein of the HIV-1 co-receptor - useful in
PT	developing resistance of CCR5-expressing cells to HIV-1 infection
PT	
XX	Claim 8; Page 31; 55pp; English.
PS	
XX	This is the nucleotide sequence of a claimed hybridisation probe
CC	used in claimed methods for detecting and identifying the presence
CC	HIV-1 CCR5 co-receptor variant nucleic acids (see AAV84125 and
CC	AAV84159) in a sample. The detection of CCR5 variants may be used to
CC	identify individuals at lower risk of infection relative to the
CC	general population who, if infected, may exhibit slower progression
CC	to AIDS.
CC	
XX	Sequence 461 BP; 102 A; 112 C; 106 G; 141 T; 0 other;
SQ	
Query Match 2.9%; Score 65; DB 20; Length 461;	
Best Local Similarity 100.0%; Pred. No. 1.4e-21;	
Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps	
QY	466 GTCCATGCTGTGGTTTGAAGGCCAGGACGGTCACCTTTGGGTGGTGACAAGTGTG 525
Dd	156 GTCCATGCTGTGGTTTGAAGGCCAGGACGGTCACCTTTGGGTGGTGACAAGTGTG 215
QY	526 ATCAC 530
Dd	216 ATCAC 220
RESULT 15	
AAT90116	
ID	AAT90116 standard; cDNA; 792 BP.
AC	
XX	AAT90116;
AC	
XX	14-APR-1998 (first entry)
DT	
XX	cDNA for inactive human CCR5.
DE	
XX	Inactive; human Cys-Cys chemokine receptor 5; CCR5;
KW	human immunodeficiency virus; type 1; type 2; HIV-1; HIV-2;
KW	predisposition; resistance; diagnosis; treatment; prevention;
KW	inflammatory disease; rheumatoid arthritis; glomerulonephritis;
KW	asthma; idiopathic pulmonary fibrosis; psoriasis; viral infection;
KW	cancer; atherosclerosis; autoimmune disorder; ss.
XX	
OS	Homo sapiens.
XX	
FH	Key Location/Qualifiers
CDS	240..792
FT	/tag= a
FT	

FT XX /note= "carboxy-terminal coding sequence deleted"
PN WO9732019-A2.
XX
PD 04-SEP-1997.
XX
PF 28-FEB-1997; 97WO-BE00023.
XX
PR 06-AUG-1996; 96EP-0870102.
PR 01-MAR-1996; 96EP-0870021.
XX
FA (EURO-) EUROSREEN SA.
XX
PI Libert F, Parmentier M, Samson M, Vassart G;
XX WPI: 1997-479829/44.
DR P-PSDB; AAW27406.
DR
XX
PT Active and inactive forms of human CC chemokine receptor CCR-5 -
PT useful to diagnose, prevent and/or treat inflammatory disorders,
PT autoimmune disease and viral infection
XX
PS Claim 18; Fig 1a; 94pp; English.
XX
CC The present sequence encodes an inactive human CC (Cys-Cys)
CC chemokine receptor 5 (CCR5), which is not a receptor of human
CC immunodeficiency virus type 1 or type 2 (HIV-1 or HIV-2). CCR5 or
CC its cDNA can be used to diagnose, treat and/or prevent inflammatory
CC diseases, e.g. rheumatoid arthritis, glomerulonephritis, asthma,
CC idiopathic pulmonary fibrosis and psoriasis, viral infections,
CC especially HIV-1 or HIV-2 infection, cancer, atherosclerosis and
CC autoimmune disorders. Subjects that express the inactive receptor
CC have a predisposition, or resistance to HIV-1 and/or HIV-2.
XX
SQ Sequence 792 BP; 208 A; 195 C; 156 G; 233 T; 0 other;

Query Match
Best Local Similarity 2.9%; Score 65; DB 18; Length 792;
Matches 65; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 466 GTCCATGCTGCTTTTAAAGCCAGGACGGTCCCTTTGGGGTGTGACAAAGTGTG 525
Db ||||||||||||||||||||||||||||||||||||||||||||||||||||
Db 630 GTCCATGCTGCTTTTAAAGCCAGGACGGTCCCTTTGGGGTGTGACAAAGTGTG 689
QY 526 ATCAC 530
Db |||||
Db 690 ATCAC 694

Search completed: June 1, 2003, 20:41:22
Job time : 1487 secs